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Smarter questions : Smarter answers

Getting A Grasp
On CRO Selection



CRO Awards



Category Winners



Company Profiles





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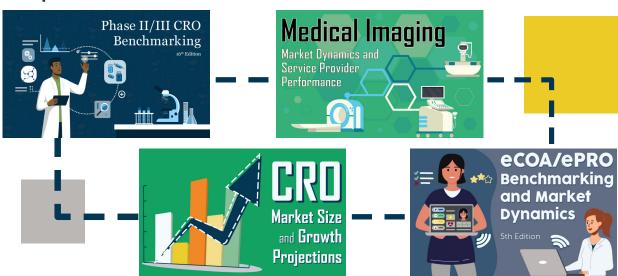




DATA-DRIVEN TOOLS FOR NAVIGATING THE CLINICAL RESEARCH SPACE

Impactful primary market research designed to offer insight into technology adoption, market dynamics, service provider selection, and service provider performance.

Recent reports include:







Getting A Grasp On CRO Selection

REBECCA MCAVOY Chief Research Officer, Industry Standard Research

The CRO selection process is a complicated one. Many factors come into play during this process, including individual trial needs, contractual arrangements, prior experience with providers, and personal preferences, just to name a few. One critical factor is undoubtedly the presence of preferred provider agreements (PPAs). Companies can spend a great deal of time negotiating these agreements, and their presence, or lack thereof, can make or break which CROs are shortlisted or awarded a Phase 2/3 study.

SR delves into service provider selection and performance in our Phase 2/3 CRO Benchmarking Study, now in its 16th year. Because preferred provider agreements can play such pivotal roles in provider selection, we tailor our research questions to ensure that they reflect the relevant decision-making scenario(s) for each respondent. First, we must understand which respondents have preferred provider agreements for Phase 2/3 services.

Among the 232 Phase 2/3 outsourcers surveyed in ISR's most recent study, over half (57%) have preferred provider agreements in place. Company size has strong ties to the usage of preferred provider agreements, with a significantly higher proportion of respondents from large sponsors (88%) reporting that they use preferred provider agreements as compared to those from midsize sponsors (54%) and small sponsors (16%).

ISR then asks respondents to share their top CRO selection criteria in a manner that reflects the roles preferred provider agreements can play in the decision-making process. Respondents whose companies have preferred providers are asked about the importance of selection criteria in two different scenarios: first, choosing from among their preferred providers and, second, choosing a provider that is not on their preferred provider list. Respondents whose companies do not use PPAs are asked about their selection criteria in a general fashion. The Venn diagram in Figure 2 is used to portray the similarities and differences in the importance of selection criteria across these three scenarios. The top seven most important attributes in each scenario are shown.

The attributes in the center of the Venn diagram represent the selection criteria respondents deemed among the most important across all three scenarios: choosing from among preferred providers, choosing a provider not on a preferred list, or choosing with no preferred provider list. "Operational excellence" has been a critical and all-encompassing attribute for project delivery over the years. "Therapeutic expertise" is

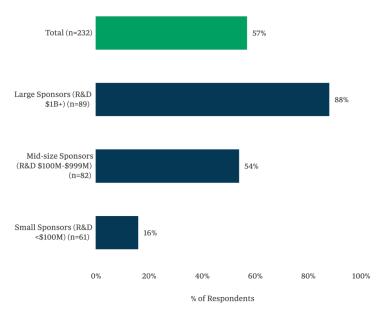


Figure 1: "Does your company have formal preferred provider agreements for Phase 2/3 services?" (n=232)

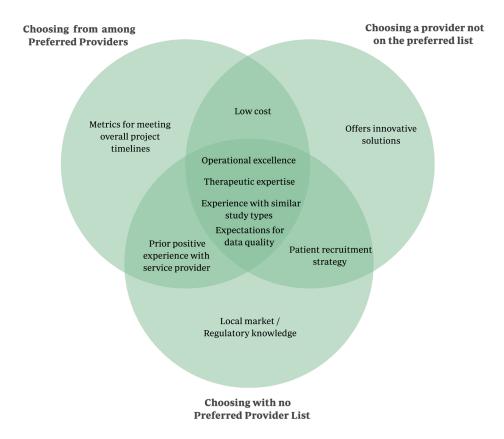


Figure 2

valued regardless of the decision-making scenario, as a thorough understanding of the therapeutic area is key to effective trial execution. "Experience with similar study types" is also important across the board, suggesting that CROs should provide examples of success with similar trials. Finally, "Expectations for data quality" has become more important recently, landing in the center of the Venn for the past three years.

Some interesting findings emerge when examining the differences among the three scenarios. For example, respondents whose companies do not have preferred providers place more importance on "Local market/Regulatory knowledge" than do respondents whose companies have preferred providers. This may be tied to the fact that respondents without PPAs are more likely to work at small companies that may not have as much in-house expertise with regulatory requirements across multiple geographies.

Another noteworthy nuance is that "Offers innovative solutions" appears as a uniquely important factor for those who are deviating from their preferred provider list. Some trials may require specialized approaches or knowledge that necessitate searching beyond the capabilities of their preferred providers.

All of this analysis now begs the question of how to put it to use. Understanding CRO selection criteria is essential for both biopharma outsourcers and CROs. Decision makers at sponsor organizations can use the insights of their peers to enhance their own approaches to Phase 2/3 outsourcing. Leadership and business development staff at CROs can use this information to better tailor their outreach and messages to target sponsors more effectively. To make these data even more powerful, coupling a grasp of sponsor selection priorities with provider performance metrics can lead to even more confident CRO selections for sponsors and marketing for CROs.



REBECCA MCAVOY is chief research officer, Industry Standard Research.

Survey Methodology: Industry Standard Research is a full-service market research provider to the pharma and pharma services industries. ISR's CRO Quality Benchmarking research is conducted annually via an online survey. For the 2024 CRO Awards data, 42 service providers were evaluated on 20+ different performance metrics. Research participants were recruited from biopharmaceutical companies of all sizes and are screened for decision-making influence and authority when it comes to working with CROs. Respondents only evaluate companies with which they have worked on an outsourced project within the past 18 months. This level of qualification ensures that quality ratings come from actual involvement with a business and that companies identified as leaders are backed by experiential data.

For more information, please visit www.ISR reports.com. 2024 CRO LEADERSHIP AWARDS WINNERS

The pharma and biopharma executives in *Clinical Leader's* and *Life Science Connect's* readership have told us about their struggles in efficiently vetting potential CRO partners. In response to this input, the CRO Leadership Awards were developed.

Based on research from Industry Standard Research's Contract Research Organization Quality Benchmarking annual online survey, 42 contract research organizations were evaluated on more than 20 different performance metrics. Research participants were recruited from biopharmaceutical companies of all sizes and were screened for decision-making influence and authority when it comes to working with contract research organizations. Respondents only evaluated companies with which they have worked on an outsourced project in the past 18 months. This level of qualification ensures that quality ratings come from actual involvement with a business and that companies identified as leaders are backed by experiential data. CROs have an opportunity to win these awards in up to three groups of outsourcing respondents — Big Pharma, Small Pharma, and Overall (combined Big and Small Pharma).

We also recognize those companies that scored one standard deviation or more above the weighted average in each of the core categories. You will see these companies noted as the *Champions*.

WHAT ARE THE AWARDS?

ISR's survey participants were asked to provide an expectation rating for each CRO they have worked with in the past 18 months. Points were then totaled for a combined score for each attribute, and a composite score for each core category was determined. Winning CROs were identified when comparing their overall score vs. the competitive set.

To learn more about ISR's industry reports, and customized research, or to be included in future CRO Leadership Awards annual surveys, visit isrreports.com or contact ISR at (919) 301-0106.

PRESENTED BY:





RESEARCH CONDUCTED BY:



Smarter questions : Smarter answers



- Access to patient populations
- Access to "unique" tests, machines, equipment
- Biostatistics
- Central lab
- Data management
- Investigator recruitment
- Monitoring
- Patient/volunteer recruitment (Phase 1)
- Patient recruitment (Phase 2/3)
- Speed of site start-up
- Technology for real-time access to data

CAPABILITIES



CHAMPIONS

OVERALL

ClinChoice

Caidya

EXCEEDED CUSTOMER EXPECTATIONS

OVERALL

CHDR

Lambda Therapeutic Research

ProPharma

Duke Clinical Research Institute

Precision for Medicine

Celerion

SGS Health Science

The PPD clinical research

business of

Thermo Fisher Scientific

ICON

Fortrea

IQVIA

Worldwide Clinical Trials

Parexel

BIG PHARMA

Precision for Medicine

Worldwide Clinical Trials

Medpace

Fortrea

ICON

SMALL PHARMA

Lambda Therapeutic Research

CHDR

Celerion

Innovaderm Research Inc.

The PPD clinical research

business of

Thermo Fisher Scientific

Caidya

IQVIA

Eurofins BioPharma Services

Novum

Altasciences

Precision for Medicine

Parexel

ICON

The terms "Small Pharma" and "Big Pharma" pertain to the outsourcing respondents, not the winners. "Overall" is a combination of Big and Small Pharma. To be recognized as a Champion, company must have scored one standard deviation or more above the weighted average in the particular category respondent group.



- Easy to work with
- Responsiveness
- Timely project communications

COMPATIBILITY



CHAMPIONS

OVERALL

CHDR

SGS Health Science

Caidya

BIG PHARMA

SGS Health Science Worldwide Clinical Trials

SMALL PHARMA

CHDR

EXCEEDED CUSTOMER EXPECTATIONS

OVERALL

ClinChoice

ProPharma

Worldwide Clinical Trials

Allucent

The PPD clinical research

business of

Thermo Fisher Scientific

Lambda Therapeutic Research

ICON

Precision for Medicine

PSI CRO AG

IQVIA

BIG PHARMA

Medpace

Precision for Medicine

Syneos Health

ICON

SMALL PHARMA

Caidya

The PPD clinical research

business of

Thermo Fisher Scientific

Allucent

Worldwide Clinical Trials

Lambda Therapeutic Research

Precision for Medicine

IQVIA

ICON

ProPharma

The terms "Small Pharma" and "Big Pharma" pertain to the outsourcing respondents, not the winners. "Overall" is a combination of Big and Small Pharma. To be recognized as a Champion, company must have scored one standard deviation or more above the weighted average in the particular category respondent group.



- Local market/regulatory knowledge
- Operational excellence
- Scientific knowledge of the Phase 1 unit's lead investigator
- Study design expertise
- Therapeutic expertise

EXPERTISE



CHAMPIONS

OVERALL

Innovaderm Research Inc.

CHDR

ClinChoice

BIG PHARMA

Worldwide Clinical Trials

SMALL PHARMA

Innovaderm Research Inc.
CHDR

EXCEEDED CUSTOMER EXPECTATIONS

OVERALL

Duke Clinical Research Institute

Caidya

IQVIA

Worldwide Clinical Trials

Allucent

SGS Health Science

Novum

PSI CRO AG

ICON

ProPharma

The PPD clinical research

business of

Thermo Fisher Scientific

Parexel

BIG PHARMA

SGS Health Science

Precision for Medicine

Fortrea

Medpace

ICON

IQVIA

SMALL PHARMA

IQVIA

The PPD clinical research

business of

Thermo Fisher Scientific

Novum

Parexel

Allucent

ICON

Worldwide Clinical Trials



- Data quality
- Project manager quality

QUALITY



CHAMPIONS

OVERALL

CHDR

ClinChoice

Caidya

BIG PHARMA

Worldwide Clinical Trials

SMALL PHARMA

CHDR

EXCEEDED CUSTOMER EXPECTATIONS

OVERALL

Duke Clinical Research Institute

Worldwide Clinical Trials

SGS Health Science

ProPharma

Precision for Medicine

Celerion

ICON

The PPD clinical research

business of

Thermo Fisher Scientific

PSI CRO AG

Fortrea

Lambda Therapeutic Research

IQVIA

BIG PHARMA

SGS Health Science

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Fortrea

ICON

SMALL PHARMA

Celerion

Caidya

The PPD clinical research

business of

Thermo Fisher Scientific

Worldwide Clinical Trials

ICON

Precision for Medicine

Lambda Therapeutic Research

IOVIA

Innovaderm Research Inc.

Quotient Sciences

The terms "Small Pharma" and "Big Pharma" pertain to the outsourcing respondents, not the winners. "Overall" is a combination of Big and Small Pharma. To be recognized as a Champion, company must have scored one standard deviation or more above the weighted average in the particular category respondent group.



- Meeting overall project timelines
- Operational excellence
- Staff turnover

RELIABILITY



CHAMPIONS

OVERALL

Duke Clinical Research Institute ClinChoice

CHDR

BIG PHARMA

Worldwide Clinical Trials

SMALL PHARMA

CHDR

EXCEEDED CUSTOMER EXPECTATIONS

OVERALL

ProPharma

Caidya

Novum

SGS Health Science

Precision for Medicine

Celerion

The PPD clinical research

business of

Thermo Fisher Scientific

Worldwide Clinical Trials

ICON

PSI CRO AG

Medpace

IQVIA

Parexel

BIG PHARMA

Precision for Medicine

Medpace

SGS Health Science

Fortrea

ICON

SMALL PHARMA

The PPD clinical research business of

Thermo Fisher Scientific

Novum

IQVIA

Innovaderm Research Inc.

ProPharma

Precision for Medicine

Caidya

Celerion

Quotient Sciences



The right outsourcing decision will help your program thrive



The inSeption Difference

The Right Culture

Attracts a subset of high-performing people who are driven by personal accountability and excellence.

Continuity of Resources

Teams value your assets as you do and are committed to seeing your program through to completion.

Predictable Costs

Our high-quality of work and unique pricing model eliminates change orders and unexpected costs.

<6%

Employee attrition vs 25-30% industry average **97%**

Customer satisfaction vs 55% reported by competitors 94%

Referral-based business from existing relationships

Put your program in trusted hands.





info@inseptiongroup.com inSeptionGroup.com



INDIVIDUAL ATTRIBUTE AWARDS

The Individual Attribute Awards were developed because of many conversations we have had with the readers of *Clinical Leader*. These conversations uncovered common attributes that sponsor companies identified as being imperative when choosing a supplier and deciding to continue doing business with a supplier.

They were often referred to as the ever-important "intangibles" a supplier brings to the table. Outside of the cover metrics of capabilities, compatibility, expertise, quality, and reliability, these attributes were what our readers identified as being most important, and as such, we felt it was important to share the data with other sponsor companies.

DATA QUALITY

TOP PERFORMERS

CHDR

Duke Clinical Research Institute

Caidya

ClinChoice

EXCEEDED CUSTOMER EXPECTATIONS

SGS Health Science

ProPharma

Lambda Therapeutic Research

Precision for Medicine

ICON

Fortrea

The PPD clinical research business of Thermo Fisher Scientific

IQVIA

MEETING PROJECT TIMELINES

TOP PERFORMERS

ClinChoice

Duke Clinical Research Institute

EXCEEDED CUSTOMER EXPECTATIONS

SGS Health Science

Innovaderm Research Inc.

ProPharma

Caidya

CHDR

Lambda Therapeutic Research

Quotient Sciences

Celerion

Worldwide Clinical Trials

ICON

The PPD clinical research business of Thermo Fisher Scientific

Precision for Medicine

Fortrea

OPERATIONAL EXCELLENCE

TOP PERFORMERS

CHDR

ClinChoice

Novum

EXCEEDED CUSTOMER EXPECTATIONS

PSI CRO AG

Worldwide Clinical Trials

Quotient Sciences

ICON

The PPD clinical research business of Thermo Fisher Scientific

IQVIA

Precision for Medicine

Caidya

Duke Clinical Research Institute

Medpace

ProPharma

Celerion



INDIVIDUAL ATTRIBUTE AWARDS

TECHNOLOGY FOR ACCESS TO DATA

TOP PERFORMERS

ClinChoice

CHDR

EXCEEDED CUSTOMER EXPECTATIONS

Duke Clinical Research Institute

Precision for Medicine

Caidya

IQVIA

SGS Health Science

Worldwide Clinical Trials

ProPharma

Parexel

The PPD clinical research business of Thermo Fisher Scientific

ICON

UBC

Novum

THERAPEUTIC EXPERTISE

TOP PERFORMERS

Duke Clinical Research Institute

ClinChoice

Innovaderm Research Inc.

EXCEEDED CUSTOMER EXPECTATIONS

PSI CRO AG

CHDR

Caidya

Novum

Allucent

Worldwide Clinical Trials

IQVIA

Parexel

Precision for Medicine

ICON

Celerion

Medpace

Eurofins BioPharma Services

Allucent caldya



CATEGORIES WON:



Allucent

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KEY LOCATIONS: Cary, NC, USA; Bracknell, Berkshire, UK; Schiphol, Amsterdam,

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CATEGORIES WON:

Caidya

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KEY LOCATIONS: Sydney, Australia; São Paolo, Brazil; Shanghai, China; Eschborn, Germany; Krakow, Poland; Seoul, South Korea; Madrid, Spain; Buckinghamshire, **United Kingdom**

CATEGORIES WON:



Celerion

Lincoln, NE, USA celerion.com

Phone: (402) 476-2811 Contact: David Maya

Email: david.maya@celerion.com

KEY LOCATIONS: Lincoln, NE, USA; Tempe, AZ, USA; West Conshohocken, PA, USA; Montreal, QC, Canada; Belfast, Northern Ireland, UK; Zürich, Switzerland; Vienna, Austria

DRUG LIFE CYCLE STAGES:

Research & Development: Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS:

Pre-Clinical, Full-Service Clinical

SERVICES & CAPABILITIES: Allucent is a global provider of comprehensive drug development solutions, including consulting, clinical operations, biometrics, and clinical pharmacology in a wide range of therapeutic areas.

THERAPEUTIC AREAS: Allucent's therapeutic areas include: Oncology & Hematology, Neuroscience, Infectious Disease & Vaccine, Immunology & Inflammation, Allergy & Asthma, Cell & Gene Therapy, Pediatrics, Rare Diseases

INDIVIDUAL ATTRIBUTE AWARDS:

Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS:

Full-Service Clinical

SERVICES & CAPABILITIES: Caidya offers fit-for-purpose solutions across early-phase development & clinical pharmacology, clinical operations, clinical science, medical affairs, medical writing, regulatory affairs & strategy, data management, biostats & programming, drug safety & pharmacovigilance.

THERAPEUTIC AREAS: Oncology, Hematology, Immuno-oncology, Cell & Gene Therapy, Rare Disease, Neurology, Nephrology, Cardiology, Ophthalmology, Dermatology, Infectious Disease, Gastroenterology, Respiratory, Women's Health

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines,

Operational Excellence, Technology for Real-Time Access to Data, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2)

MAIN SERVICE AREAS:

Lab, Full-Service Clinical

SERVICES & CAPABILITIES: Celerion excels at Phase 1-2 clinical pharmacology ranging from PK/ PD, FIH, SAD/MAD, DDI, QTc, Bioequivalence, and ADME studies to multisite patient studies with full services (project management, data management, statistics, regulatory support & bioanalytical services).

THERAPEUTIC AREAS: Cardiovascular Disease, Renal Insufficiency, Hepatic Insufficiency, Immunotherapy, Infectious Disease, Metabolic Disease, Oncology, Respiratory Disease, Vaccines, and others

INDIVIDUAL ATTRIBUTE AWARDS: Meeting Overall Project Timelines, Operational Excellence, Therapeutic Expertise









Duke Clinical Research Institute

Durham, NC, USA dcri.org

Phone: (919) 668-8700 Contact: Carolyn Arias

Email: carolyn.arias@duke.edu

KEY LOCATIONS: Durham, NC

CATEGORIES WON:



Eurofins BioPharma Services

Lancaster, PA, USA eurofins.com/biopharma-services/

Email: clinicaltrials@bcl.eurofins.com

KEY LOCATIONS: The Eurofins BioPharma Services Network of Companies has a global footprint in the USA, Europe, and APAC regions.

CATEGORIES WON:

Fortrea

Durham, NC, USA fortrea.com

Email: GCOInsideSalesLeadership@fortrea.com

KEY LOCATIONS: Durham, NC, USA; Dallas, TX, USA; São Paulo, Brazil; Leeds, UK; Munich, Germany; Shanghai, China; Bangalore, India; Tokyo, Japan

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS:

Full-Service Clinical

SERVICES & CAPABILITIES: Our services include tailored trial design, collaborative academic leadership, established NA investigator networks, efficient data collection, risk-based monitoring, and shared endpoint adjudication.

THERAPEUTIC AREAS: Renowned for cardiology, we pioneer trials in GI health, infectious disease, neuroscience, oncology, pediatrics, and respiratory medicine. Our depth of knowledge is shared via impactful publications and global meetings.

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines, Operational Excellence, Technology for Real-Time Access to Data, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

Drug Substance Production: Primary Process Development, Drug Substance Production

Formulated Drug Production: Dosage Form Development, Dosage Form Production, Packaging, Logistics

MAIN SERVICE AREAS:

Lah

SERVICES & CAPABILITIES: The Eurofins BioPharma Services Network of Companies is a first-class biopharmaceutical outsourcing services partner. The network covers the whole chain of drug development with a global footprint, uniform QA systems, and high-quality services.

THERAPEUTIC AREAS: Oncology, Cell & Gene Therapy, Vaccine Development, Cardiovacular Disease, Immunology, CNS, Gastroenterology, Pulmonary/Respiratory Diseases

INDIVIDUAL ATTRIBUTE AWARDS:

Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS:

Full-Service Clinical

SERVICES & CAPABILITIES: Integrated clinical pharmacology solutions, Phase 1 through Phase 4 full-service clinical development, FSP and hybrid models, diagnostic and device development, regulatory strategy, development, market access, and HEOR consulting

THERAPEUTIC AREAS: Cardiovascular and metabolism; cell, gene, and advanced therapies; hepatology; immunology and inflammation; infectious diseases and vaccines; nephrology; neurology and ophthalmology; oncology; rare disease and pediatrics; respiratory; metabolic diseases; rare disease; and immunology

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines









ICON

Dublin, Ireland iconplc.com

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Email: Lisa.King@iconplc.com

KEY LOCATIONS: Dublin, Ireland; Paris, France; Reading, United Kingdom; Blue Bell, PA, USA; Raleigh, NC, USA; Tokyo, Japan; Sydney, Australia; São Paulo, Brazil



CATEGORIES WON: Innovaderm Research Inc.

Montreal, QC, Canada innovaderm.com/full-service-dermatology-cro

Phone: 1-866-575-3111 Contact: Audrey Miron

Email: busdev@innovaderm.com

KEY LOCATIONS: Barcelona, Spain; Warsaw, Poland; California, USA; Florida, USA; Boston, MA, USA; New York, USA; Edmonton, AB, Canada; Toronto, ON, Canada; Montreal, QC, Canada

CATEGORIES WON:

IQVIA

Durham, NC, USA

iqvia.com

Contact: Jonathan Limbouris Email: jonathan.limbouris@igvia.com

KEY LOCATIONS: Global

DRUG LIFE CYCLE STAGES:

Research & Development: Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

Formulated Drug Production: Dosage Form Development, Dosage Form Production, Packaging, Logistics

MAIN SERVICE AREAS: Lab, Full-Service Clinical

SERVICES & CAPABILITIES: ICON offers a full range of clinical, consulting, and commercial services from clinical development strategy, planning, and trial design to full study execution and post-market commercialization.

THERAPEUTIC AREAS: Oncology, Central Nervous Systems, Infectious Diseases and Vaccines, Cell and Gene Therapies, Cardiovascular, Endocrine and Metabolic, Ophthalmology, Paediatrics, Women's Health, Internal Medicine, and Immunology

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines, Operational Excellence, Technology for Real-Time Access to Data, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS: Full-Service Clinical

SERVICES & CAPABILITIES: Innovaderm is THE leading full-service dermatology CRO built upon deep dermatology expertise and proven success, offering full operational capabilities for all phases of clinical development toward the benefit of improving patients' lives.

THERAPEUTIC AREAS: 30+ dermatology indications including: Acne, Alopecia Areata, Atopic Dermatitis, Hidradenitis Suppurativa, Notalgia Paresthetica, Psoriasis, PPP, GPP, Rosacea, and more. Dermal Fillers & other Medical Aesthetics trials. Systemic Sclerosis & other Rheumatology indications.

INDIVIDUAL ATTRIBUTE AWARDS: Meeting Overall Project Timelines, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3)

Drug Substance Production: Drug Substance Production

MAIN SERVICE AREAS: Lab, Full-Service Clinical

SERVICES & CAPABILITIES: IQVIA offers full service and functional service models to support pharma companies in Phase I-3 research, including patient recruitment and retention, clinical monitoring, life cycle safety, laboratory services, data management, biostatistics, and medical writing.

THERAPEUTIC AREAS: Oncology, CNS, Infectious Disease & Vaccines, Cardiovascular, Dermatology, GI & Hepatology, Endocrinology, Allergy & Respiratory, Rheumatology, Ophthalmology, Nephrology, & Reproductive Health

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Operational Excellence, Technology for Real-Time Access to Data, Therapeutic Expertise

MEDPRCE PO(EXE propharma



CATEGORIES WON:



Medpace

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KEY LOCATIONS: Cincinnati, OH, USA; Shanghai, China; Singapore; Tokyo, Japan; Leuven, Belgium; London, UK; München,

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CATEGORIES WON:

Parexel

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China

CATEGORIES WON:



Raleigh, NC, USA propharmagroup.com

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KEY LOCATIONS: Raleigh, NC, USA; Chicago, IL, USA; London, UK; Leiden, Netherlands; Lund, Sweden; Berlin, Germany. Regions: Central Eastern Europe;

Australia; Japan; China; India

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2, Phase 3)

MAIN SERVICE AREAS: Full-Service Clinical

SERVICES & CAPABILITIES: A full range of integrated services: Medical Affairs, Regulatory, Clinical Monitoring, Clinical Trial Management, Biometrics, Safety & Pharmacovigilance, Quality Assurance, Technology

THERAPEUTIC AREAS: Autoimmune Diseases, Cardiovascular, Cellular & Gene Therapies, Dermatology, Endocrine & Metabolic, Gastroenterology, Hematology & Oncology, Infectious Diseases & Vaccines, Hepatology, Nephrology, Neuroscience, Ophthalmology, Pediatrics, Radiation Therapy, Rare Disease & Orphan Indications, Women's Health

INDIVIDUAL ATTRIBUTE AWARDS:

Operational Excellence, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 1, Phase 2. Phase 3. Phase 4)

Formulated Drug Production: Packaging, Logistics

MAIN SERVICE AREAS: Lab, Full-Service Clinical

SERVICES & CAPABILITIES: From clinical trials to regulatory and consulting to commercial and market access, our therapeutic, technical, and functional ability and services help life sciences and biopharmaceutical clients transform their scientific discoveries into treatments.

THERAPEUTIC AREAS: Oncology, hematology, infectious disease & vaccines, neurology, respiratory, dermatology, endocrine & metabolism, rheumatology, cardiovascular, gastroenterology, genitourinary & nephrology, psychiatry, ophthalmology, allergy & immunology, women's health, and orthopedics

INDIVIDUAL ATTRIBUTE AWARDS: Technology for Real-Time Access to Data, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

Drug Substance Production: Primary Process Development

MAIN SERVICE AREAS: Pre-Clinical, Full-Service Clinical

SERVICES & CAPABILITIES: ProPharma's deep domain expertise in regulatory affairs, clinical research, quality & compliance, pharmacovigilance, medical information, & R&D tech offers an end-to-end suite of fully customizable solutions that de-risk & accelerate drug & device programs for sponsors.

THERAPEUTIC AREAS: ProPharma offers custom & proven solutions across the following therapeutic areas: Oncology, Rare Disease, Cell & Gene Therapy, Cardiovascular, Cell & Gene Therapy, Central Nervous System (CNS), Infectious Disease/Vaccines, Ophthalmology, Pediatrics, Respiratory System, Blood Diseases, and Immune System.

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines, Operational Excellence, Technology for Real-Time Access to Data



PSI CRO AG

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Email: olga.alfonsova@psi-cro.com

KEY LOCATIONS: Zug, Switzerland; Philadelphia, PA, USA; San Francisco, CA, USA; Warsaw, Poland; Munich, Germany; Buenos Aires, Argentina; Sydney, Australia; Seoul, South Korea



Molecule to cure. Fast.™

CATEGORIES WON:

Ouotient Sciences

Nottingham, UK quotientsciences.com

Email: info@quotientsciences.com

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CATEGORIES WON:

SGS Health Science

Geneva, Switzerland sgs.com/healthscience

Phone: +41 22 739 91 11 Contact: SGS Health Science Email: healthscience@sgs.com

KEY LOCATIONS: Geneva, Switzerland; Antwerp, Belgium; Mississauga, ON, Canada; Lincolnshire, IL, USA; Berlin, Germany; Glasgow, Dumbartonshire, UK; Shanghai, China; Paris, France

DRUG LIFE CYCLE STAGES:

Research & Development: Clinical (Phase 2, Phase 3)

MAIN SERVICE AREAS: Full-Service Clinical

SERVICES & CAPABILITIES: PSI CRO is a privately owned, full-service CRO operating globally. PSI's global reach supports clinical trials across multiple countries and continents and specializes in the planning and execution of global pivotal registration clinical trials.

THERAPEUTIC AREAS: Oncology, including breast cancer, lung cancer, prostate cancer, colorectal cancer, ovarian cancer, and radiopharmaceuticals; hematology-oncology and hemophilia; gastrointestinal diseases, including Crohn's disease, ulcerative colitis, and other inflammatory bowel diseases

INDIVIDUAL ATTRIBUTE AWARDS:

Operational Excellence, Therapeutic Expertise

DRUG LIFE CYCLE STAGES:

Research & Development: Pre-Clinical, Clinical (Phase 1, Phase 2)

Drug Substance Production: Primary Process Development, Drug Substance Production

Formulated Drug Production: Dosage Form Development, Dosage Form Production, Packaging, Logistics

MAIN SERVICE AREAS: Lab, Pre-Clinical, **Full-Service Clinical**

SERVICES & CAPABILITIES: Drug substance (API) synthesis & development, API manufacturing, drug product formulation development, drug product manufacturing, Phase 1 clinical trials with healthy volunteers, bioanalysis, human ADME, 14C radiolabeled drug substance, data sciences, and clinical pharmacology

THERAPEUTIC AREAS: Orphan Drugs for Rare Diseases, Oncology, Pediatrics, Cardiology, Gastroenterology/GI, Pain Management, CNS & Neurology, Psychedelic Therapeutics

INDIVIDUAL ATTRIBUTE AWARDS: Meeting Overall Project Timelines, Operational Excellence

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3)

Drug Substance Production: Primary Process Development

Formulated Drug Production: Dosage Form Development, Dosage Form Production, Packaging, Logistics

MAIN SERVICE AREAS: Lab, Pre-Clinical, **Full-Service Clinical**

SERVICES & CAPABILITIES: Formulation development, quality control testing for raw materials, APIs, and finished products; manufacturing for Phase 1-3 clinical trials: full clinical research services with an SGS clinical pharmacology unit, central lab services, and bioanalysis

THERAPEUTIC AREAS: SGS supports all therapeutics areas with formulation, QC testing, clinical manufacturing and clinical research. SGS clinical research services specialize in infectious disease, respiratory, and CNS.

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines, Technology for Real-Time Access to Data





Syneos Health

Morrisville, NC, USA syneoshealth.com

Email: meg.byers@syneoshealth.com

KEY LOCATIONS: Syneos Health has offices in North America, Latin America, Europe, Asia Pacific, the Middle East, and Africa.



CATEGORIES WON:



The PPD clinical research business of Thermo Fisher Scientific

Wilmington, NC, USA thermofisher.com/ppd

Email: salesops@ppd.com

KEY LOCATIONS: Wilmington, NC, USA; Research Triangle Park, NC, USA; Middleton, WI, USA; Richmond, VA, USA; Austin, TX, USA; China; Japan; Asia-Pacific; UK; Europe; Middle East; Africa; Latin America



CATEGORIES WON ;



Worldwide Clinical Trials

Research Triangle Park, NC, USA worldwide.com

Email: OfficeofCEO@worldwide.com

KEY LOCATIONS: Research Triangle Park, NC, USA; Austin, TX, USA; Boston, MA, USA; Warsaw, Poland; Belgrade, Serbia; Nottingham, UK; São Paulo, Brazil; Tokyo, Japan

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS: Full-Service Clinical

SERVICES & CAPABILITIES: Syneos Health® is a leading fully integrated biopharmaceutical solutions organization built to accelerate customer success. We translate unique clinical, medical affairs, and commercial insights into outcomes to address modern market realities.

THERAPEUTIC AREAS: Biosimilars, Cardiovascular, Cell & Gene Therapy, Neuroscience, Dermatology, Endocrine & Metabolic, Gastroenterology, Immunology & Inflammation, Infectious Diseases, Oncology & Hematology, Pediatrics, Rare Disease, Respiratory, Women's Health, and more

DRUG LIFE CYCLE STAGES:

Research & Development: Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

MAIN SERVICE AREAS: Lab, Full-Service Clinical

SERVICES & CAPABILITIES: Phase 1-3b clinical development & laboratory services: bioanalytical, biomarkers, central, GMP, vaccine sciences; AI/ML; biotech; digital/decentralized trials; FSP; peri- and post-approval; RWE/RWD; site/patient services; vaccine development

THERAPEUTIC AREAS: Cardiovascular, Cell/ Gene Therapy, Critical Care, Dermatology, Gastrointestinal, Immunology, Infectious Diseases, Metabolic/Endocrine, Neuroscience, Oncology/ Hematology, Pediatrics, Rare Diseases, Respiratory, Urology, Vaccines, Women's Health, and more

INDIVIDUAL ATTRIBUTE AWARDS: Data Quality, Meeting Overall Project Timelines, Operational Excellence, Technology for Real-Time Access to Data

DRUG LIFE CYCLE STAGES:

Research & Development: Discovery, Pre-Clinical, Clinical (Phase 1, Phase 2, Phase 3, Phase 4)

Formulated Drug Production: Logistics

MAIN SERVICE AREAS: Lab, Pre-Clinical, Full-Service Clinical

SERVICES & CAPABILITIES: Bioanalytical laboratory services, Phase 1-4 clinical trials, and post-approval and real-world evidence studies - all powered by an accessible team of clinicians, scientists, and researchers who bring a collaborative, personalized approach to each clinical program.

THERAPEUTIC AREAS: Neuroscience, Oncology, Rare Disease, and Cardiometabolic and Inflammatory Disease

INDIVIDUAL ATTRIBUTE AWARDS: Meeting Overall Project Timelines, Operational Excellence, Technology for Real-Time Access to Data, Therapeutic Expertise





Laval, QC, Canada altasciences.com

Email: contact@altasciences.com

KEY LOCATIONS: Laval, QC, Canada; Montréal, QC, Canada; Kansas City, KS, USA; Seattle, WA, USA; Philadelphia, PA, USA; Los Angeles, CA, USA; Scranton, PA, USA; Columbia, MO, USA; Sacramento, CA, USA

DRUG LIFE CYCLE STAGES:

Research & Development: Pre-Clinical, Clinical (Phase 1, Phase 2)

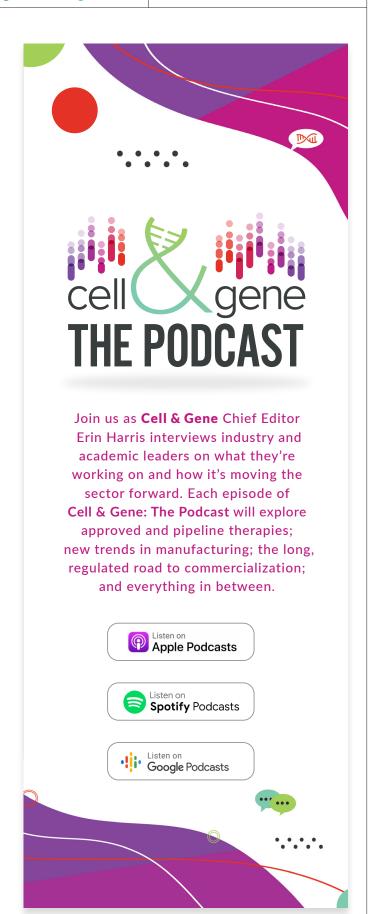
Formulated Drug Production: Dosage Form Development, Dosage Form Production

MAIN SERVICE AREAS:

Lab, Pre-Clinical, Full-Service Clinical

SERVICES & CAPABILITIES: Altasciences is an integrated CRO/CDMO offering a flexible approach to pre-clinical and clinical pharmacology studies, including bioanalysis, formulation, manufacturing, and analytical services. Other services include program management, medical writing, and biostatistics.

THERAPEUTIC AREAS: Therapeutic areas include (but are not limited to) the following: cardiology, dermatology, gastroenterology, immunology, CNS/neurology, oncology, ophthalmology, substance abuse, psychiatry, rheumatology, urology, infectious diseases, and metabolism and endocrinology.



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